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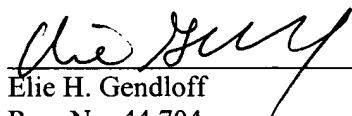


IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

St. Louis, Missouri
October 30, 2000

CERTIFICATE OF MAILING

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In re application of:
Curtiss and Tinge

Serial No.: 08/473,789

Filed: June 7, 1995

For: Recombinant Bacterial Vaccine System
With Environmentally Limited Viability

Examiner V. Ryan

Group Art Unit 1641

BOX NON FEE AMENDMENT
Assistant Commissioner for Patents
Washington, D.C. 20231

PROPOSED RESPONSE TO OFFICE ACTION UNDER 37 C.F.R. §1.111

This paper is submitted as a proposed response to the Office Action of July 25, 2000.

Applicants request that the examiner have a telephone interview with the undersigned attorney to resolve any remaining issues in this case. Applicants propose that the following remarks be entered in this case, subject to the results of the proposed telephone interview. Claims 1-4, 8-14, 16, 20, 23, 24, 27-32, 35, 37 and 41-45 are pending in this case.

The withdrawal of all previous claim rejections is noted with appreciation. Discussion regarding the current rejections is provided below.

Rejections under the non-statutory doctrine of obviousness-type double patenting

Claims 1-4, 8-14, 16, 20, 23, 24, 27-32, 35, 37 and 41-44 are rejected under the judicially created doctrine of obviousness-type double patenting over U.S. Patent Application Ser. No. 08/761,769. As there are no allowable claims in the instant application at this time, applicants wish to defer responding to this rejection until such time as there are allowable claims.

Rejections under 35 U.S.C. §102(e)

Claims 1-4, 8-14, 20, 23, 24, 27-29 and 37 are rejected under 35 U.S.C. §102(e) as being anticipated by Molin et al., U.S. Patent No. 5,702,916. It is asserted that Molin et al. discloses a biological containment system comprising a cell containing a recombinant DNA molecule that regulatably expresses a cell-killing function in certain environmental conditions. Specifically, the *hok* gene is cited as expressing a cell-killing function when insufficient concentrations of inhibitory *sok* are present.

Applicants respectfully request reconsideration and withdrawal of this rejection in light of the following remarks.

Applicants first note that all of the rejected claims contain the limitation that the essential gene is a copy of a native gene, wherein the native gene is inactivated in the cell. This limitation is not taught by the cited Molin et al. patent. Specifically, the Office Action identifies the *sok* gene of Molin et al. as an essential gene within the context of the instant invention. Applicants contend that this gene is not a native gene, nor is an inactivated copy of that gene present in the cell, as is required by the rejected claims. In Molin, the *sok* gene is carried on the plasmid R1, in the *parB* region (Molin et al., column 10, lines 24-49). Molin does not teach that the plasmid-borne *sok* is a copy of a native gene, nor does Molin teach the presence of an inactivated *sok* in the cell.

In addition, all of the rejected claims require that the essential gene be expressed when the cell is in a permissive environment, and not expressed when the cell is in a non-permissive environment. Applicants contend that this limitation is not taught by Molin et al. Specifically, the *sok* gene is not described in Molin as being differentially expressed when in permissive versus non-permissive environments. Rather, the cell killing function is differentially expressed by placing a regulatable promoter upstream of the *hok* gene such that the *hok* mRNA is expressed at a higher level than the *sok* mRNA when the cell is in a non-permissive environment (see Molin et al., column 11 lines

*Let this
essential
(native)
gene
ascl*

26-39). Molin et al. teaches a containment system which functions by regulatably controlling expression of a lethal gene, and makes no suggestion of controlling the expression of an essential gene. In all of the descriptions of the containment system in Molin et al., the *sok* gene is expressed regardless of whether the cell is in a permissive or non-permissive environment. See for example, Molin et al. at column 21, line 30 through column 23, line 60, for descriptions of the disclosed plasmids. Note, in particular, that there is no regulatable promoter controlling expression of the *sok* gene found in any of these constructs. Thus, the claim limitation that "the essential gene is expressed when the cell is in the permissive environment and is not expressed when the cell is in the non-permissive environment" is not taught by Molin et al.

The Office Action points out, at the bottom of page 6 to the top of page 7, that Molin et al. teaches a system wherein the death of the cell occurs due to the expression of the gene encoding the cell killing function as a result of pre-determined environmental conditions. In contrast, in the claimed invention of the instant application, cell death occurs both because of expression of a lethal gene in a non-permissive environment, and because of non-expression of the essential gene in non-permissive environments.

Claims 1, 4, 10-12, 20, 27, and 41-45 are rejected under 35 U.S.C. §102(b) as being anticipated by Curtiss III (*Engineering Organisms For Safety: What Is Necessary?*, The Release Of Genetically-Engineered Micro-Organisms, M. Sussman et al., editors, Academic Press, 7-20 (1988)). It is alleged in the Office Action at page 7 that Curtiss III teaches a system in which the *hok* gene is the lethal gene and *asd* is the essential gene.

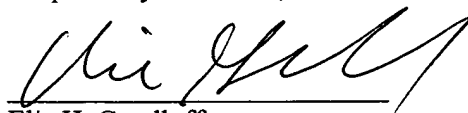
Applicants first point out that all of the rejected claims contain the limitation that "the essential gene is expressed when the cell is in the permissive environment and is not expressed when the cell is in the non-permissive environment." This limitation is not taught by Curtiss III. Specifically, the section at page 12 of Curtiss III that is emphasized by the Office Action suggests the use of a lethal gene such as *hok* in order to limit the vector of the balanced lethal system to the intended host. In that case, the essential gene, *asd*, is expressed in a non-regulatable manner, in order to provide selective pressure to maintain the vector in the host population by causing death of the host in the absence of the vector. In such a system, regulatable expression of the essential gene is not necessary or desirable. The cell killing function is suggested as a means of preventing expression of the vector by unintended hosts. The promoter suggested, therefore, is one that is repressed so as not to express the lethal gene when in the intended host cell. When the plasmid is transferred to an unintended host, however, the promoter causes expression of the lethal gene. The utility of this system does not extend to causing the death of the intended host organism when released from a permissive environment. Instead, such organisms may remain viable and proliferate in unintended environments.

In contrast, the system of the instant invention insures death of the intended host organism when the host organism is released from a specific permissive environment into a non-permissive environment, as well as death of the intended host when the vector is released, and death of unintended hosts when the vector is transferred to them. The cell-killing function of the instant invention is not only triggered by loss of the vector from the host, as it is in the Curtiss III system, but more importantly, by the release of the host organism from the permissive environment. This is accomplished in the instant invention by utilizing as the essential gene a native gene of the host organism, which has been inactivated in the host, under control of a regulatable promoter. Thus, when the host is released into the non-permissive environment, the essential gene is no longer expressed, resulting in the death of the host. In addition, death of the host organism in non-permissive environments is insured by regulatably expressing a lethal gene, such that when the organism is released from the permissive environment, the promoter is up regulated resulting in cell death by expression of the lethal gene product.

In summary, the cited references teach genetically engineered organisms that contain vectors comprising both lethal genes and essential genes. However, neither Curtiss III nor Molin et al. teaches an essential gene that is under the control of a regulatable promoter. In addition, Molin et al. does not even teach an essential gene that is a native gene, nor does it teach the presence of the inactivated native gene in the cell. Therefore, applicants assert that the claims of the instant application are not anticipated.

Based on the discussion presented above, reconsideration and withdrawal of the rejections put forth in the Office Action dated July 25, 2000 is respectfully requested. Applicants believe that the currently pending claims are allowable. If there are any issues yet to be resolved, the Examiner is invited to contact the undersigned attorney.

Respectfully submitted,



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